The listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 10 (Currently Amended) A method for regulating fertility without additional use of a follicular sex steroid comprising administering to a patient in need thereof a pharmaceutical composition comprising a 17-Chloro-D homosteroid of formula I according to claim 12.

Claim 15 (Currently Amended) A process for the production of a 17-chloro-Dhomosteroid of the formula I according to claim 12,

[paste structure]

(I)

comprising converting a 17-chloro-1,3,5(10),16-tetraene-17-one of formula II

[paste structure]

(II)

in which

 R_1 is a hydrogen atom, a $C_{1.5}$ alkyl radical, a $C_{1.6}$ alkanoyl radical or a benzoyl radical, R2 is C1-6 alkyl group,

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with a magnesium-organic reagent of general formula BrMg alkyl, BrMg alkenyl or BrMg alkinyl or with acctylene or an alkyl- or aryl-substituted acctylene in the presence of a base, or with a lithium-organic compound or with a silicon-organic compound into a 17aαsubstituted compound of formula III,

[paste structure]

(III)

in which

 R_1 is a hydrogen atom, a C_{1-6} alkyl radical, a C_{1-6} alkanoyl radical or a benzoyl radical, R_2 is a C_{1-6} alkyl group,

Ra is a hydrogen atom, a metal atom or a silvl group, and

 R_a is a hydrogen atom, a C_{1-6} alkyl group, a C_nF_{2n-1} group, in which n=1,2 or 3, or a $C=CR_3$ group, in which R_5 is a hydrogen atom, a C_{1-6} alkyl radical or an unsubstituted or substituted phenyl radical,

whereby wherein in the case of R_3 = hydrogen, the free 17a α -ethinyl compound of general formula III is further modified by a SONAGASHIRA reaction to form compounds with R_3 = $C_6H_4R_6$, in which R_6 stands for a free or substituted hydroxyl group, amino group, thiol group, sulfamate group, sulfamate group or a $C_{1.6}$ alkyl group or a $C_{6.12}$ aryl group.

Claim 17 (Currently Amended) The process according to claim 15, wherein said compound of formula $\frac{111}{2}$, in which R_1 is an acyl radical, is converted by ether cleavage into a free hydroxyl groups.

Claim 18 (Currently Amended) The process according to claim 15, wherein said

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compound of formula # III in which R3 is a hydrogen atom, is converted into others or esters.

Claim 19 (Currently Amended) A method for contraception in women comprising administering to a woman in need thereof a therapeutically effective amount of a compound of formula 1 according to claim 12.

Claim 20 (Currently Amended) A method for contraception in men comprising administering to a man in need thereof a therapeutically effective amount of a compound of formula I according to claim 12.

Claim 21 (Currently Amended) A method for treating benign or malignant proliferative diseases of the ovary comprising administering to a patient in need thereof a therapeutically effective amount of a compound of formula I according to claim 12.

Claim 25 (Currently Amended) A pharmaceutical composition according to claim 24 +2, further comprising a GnRH antagonist, a progesterone receptor antagonist, a mesoprogestin, a gestagen or a tissue-selective gestagen.